CASE REPORT

FIBRO DYSPLASIA OSSIFICANS PROGRESSIVA

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We present a case of 7 years old boy with a very rare debilitating autosomal dominant disorder characterized by heterotopic ossification. Fibro dysplasia ossificans progressiva affects 1 in 2 million individuals with only 2 previous cases reported from this region. The disease manifests as multiple foci of bone formation in muscles, fasciae, tendons and ligaments often triggered by trauma. The child was born with bilateral short hallux valgus and aplasia of distal phalanges of both thumbs. In the last 3 years he had developed hard bony swellings in the scalp, followed by limitation of neck mobility. He developed palpable nodules on the right lateral thoracic cage over the last 1 year following trauma. Heterotopic bone formation was also seen in both tibias. FOP causes irreversible lesions of ossification thus early institution of prophylactic measures, counselling regarding avoidance of trauma and surgery can significantly reduce acute exacerbations of this rare disease.

Keywords: Fibro dysplasia ossificans progressive; Heterotopic ossification; Hallux valgus

INTRODUCTION

Fibro dysplasia Ossificans Progressiva or more commonly Myositis Ossificans is a very rare autosomal dominant disorder comprising of ectopic osteogenesis in fasciae, ligaments, muscles, ligaments and joint capsules. It was first described by Guy Patin in 1968. The disease is mainly diagnosed clinically on the basis of two features, i.e., The foci of heterotopic bone formation and congenital deformities of the big toes. The prevalence rate is 1: 2,000,000. The disease is pan-ethnic and doesn't have a gender predisposition.

Overexpression of bone morphogenetic protein (BMP-4) has been found to be associated with the disease with most cases resulting from point mutations. There are approximately 800 known patients of the disease worldwide. Increased awareness of the disease is of utmost importance as accurate and early diagnosis is fundamental to prevention of progression to severe disease.

We present a case of a seven-year-old boy who presented with multiple hard bony swellings in the scalp, neck and thorax with the aim of highlighting the challenging nature of the disease and to highlight the triggers for the bone formation (trauma & surgery). Since there is no definite cure available for the disease at present therefore minimizing the acute flare-ups and prevention of formation of newer bony swellings is the prime target.

CASE REPORT

A 7-year-old school going boy was brought to outpatient clinic of Paediatric Surgery Department, Military Hospital Rawalpindi with a history of development of multiple hard bony swellings over the scalp, neck and thorax over the past 03 years. The bony swellings developed gradually secondary to traumatic falls and were now restricting the expansion of the chest on the right side. The child also had restricted mobility of neck on the right side. (Figure 3, 4) He had no other systemic complaints.

He was the first-born full-term baby boy with normal developmental milestones born of a consanguineous marriage. There was no family history of any skeletal or connective tissue disease. He was born with bilateral hallux valgus deformity and aplasia of distal phalanges of both thumbs. (Figure 1, 2) There was no other congenital anomaly in the child. He first developed hard swellings in the scalp after a fall at home at the age of 4 years but the parents did not consult any doctor for it. Over the course of next couple of years, he developed nodules on the right side of neck which were initially soft but gradually developed into hard bony swellings. The trigger for the formation of neck swellings was not known. However, neck movements on the right side were compromised. The child developed palpable hard nodules on the right lateral thoracic cage over the last 1 year following a traumatic fall while riding a bicycle. As the swellings hardened, there was restriction of his chest expansion more pronounced on the right side. He had a number of consultations from various hospitals where he was advised multiple options ranging from conservative treatment to surgery.
Figures-1 and 2: Clinical photograph showing aplasia of distal phalanges of both thumbs and bilateral hallux valgus deformity of big toes.

Figure-3: Shows clinical photograph showing multiple swellings on the right side of chest and back

Figure-4: Shows swellings on right side of neck.

Figures-5, 6: The Chest X-Ray and HRCT Chest with 3-D Reconstruction showing ectopic bone foci.
The radiograph of both tibia and fibula revealed medial tibial osteochondromas bilaterally. (Figure 9) Based on the history, examination and investigation a diagnosis of fibro dysplasia ossificans progressive was made.

DISCUSSION
Fibro dysplasia ossificans progressive (FOP) also known as Munch Meyer’s disease is a rare debilitating autosomal dominant disease characterized by progressive ectopic mature lamellar bone formation in the connective tissues of the body. Activating heterozygous mis-sense mutations in the ACVR1 gene (activin A receptor, type I) located on chromosome 2, which encodes the bone morphogenetic protein (BMP) type I receptor protein, were discovered by Shore et al in 2006 as the cause of FOP.

Most of the patients present in early childhood with development of soft tissue swellings in the scalp, neck and upper torso which gradually ossify into hard bony stigmata. The swellings may initially be painful with compromise of movements of the affected body parts. There are two characteristic features of the disease, i.e. Presence of heterotopic bone formation and hallux valgus deformity of the big toes since birth. Presence of hallux valgus deformity of big toes is pathognomonic of the condition with the deformity reported in almost 100% of the cases. Maftei et al in 2014 reported the first case of FOP that was diagnosed prenatally on ultrasound by identification of the hallux valgus deformity of the big toes. Therefore genetic testing should commence in all children born with hallux valgus deformity of big toes. More than 50% cases of the disease also have medial tibial osteochondromas (seen in our patient).

There is no gender or ethnic predisposition for FOP. The triggers for ectopic bone formation in connective tissues include blunt trauma during routine activities or surgical procedures however spontaneous onset has also been reported in literature. The mean age of onset of symptoms is generally five years. The sternocleidomastoid muscle is usually the initial site of involvement with the disease progressing down to involve the thoracic and lumbar regions. In severe cases there is bridging between axial and appendicular
skeleton, between ribs (as was the case in our patient) and between thorax and pelvis with severe restriction of motion. The respiratory complications include recurrent infections and atelectasis which usually occur due to restriction of chest expansion secondary to disease progression. The disease is misdiagnosed in up to 90% of cases and resultantly may cause severe iatrogenic injuries by doing hasty surgeries.\(^1\) The differential diagnosis includes osteosarcoma, soft tissue sarcoma, dermatomyositis, rheumatoid arthritis, desmoid tumours, aggressive juvenile fibromatosis and acquired heterotopic ossification.\(^4\) Laboratory investigations in these patients are usually normal. Plain radiographs can identify the deformities of hand and feet and can also identify bony bridges in different fascial planes. CT scan and MRI can detect the lesions in the perosseous stages which may manifest as oedema of the fascial planes and muscular bundles. Bone scan using Gallium 67 citrate and Tc-99 diphosphonate can also help to diagnose the lesions in earlier stages.\(^5\)\(^,\)\(^6\) Biopsy of lesions is avoided since it can itself be a trigger to bone formation.\(^1\)

Fibro dysplasia ossificans progressiva is one of the diseases in which molecular gene therapy (signal transduction inhibitors etc) directed at blocking Activin A type 1 receptor (ACVR1) might promise recovery from the predicament of the sufferers in future.\(^1\)\(^3\) Definitive cure is not available yet and current focus of treatment is on early diagnosis, prevention of trauma, and avoidance of invasive procedures. Different treatment regimens including steroids, bisphosphonates, NSAIDS and avo\(^1\)dance of invasive procedures. Different treatment regimens including steroids, bisphosphonates, NSAIDS, rosiglitazone, and even radiotherapy have been tried to treat the disease but the disease progression was never curtailed.\(^1\)\(^3\) In almost all instances in which surgery was performed resulted in recurrence and aggravation of the heterotopic bone formation. Steroids, NSAIDS and bisphosphonates have been effective in treating acute flare-ups, result in improvement of mobility and decrease the disease severity.\(^1\)\(^3\)\(^,\)\(^14\)

**CONCLUSION**

Since there is no curative treatment available for patients with fibro dysplasia ossificans progressive therefore the primary goal is the earliest possible diagnosis of FOP before the heterotopic ossification occurs so as to avoid trauma or any procedure or treatment that could initiate the disease process. Children with congenital deformities of hands and feet particularly hallux valgus of big toes, possibly picked on prenatal ultrasound should raise alarm. Thorough counselling of the patient and attendants regarding avoidance of trauma and invasive procedures and early institution of prophylactic and precautionary measures, can significantly reduce the acute exacerbations of this rare disease. Steroids, NSAIDS and bisphosphonates are the mainstay of treatment during acute flare-ups. Gene therapy directed at ACVR1 may give hope to the patients in future.

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**Informed consent:** Informed consent was obtained from patient's father.

**REFERENCES**


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